Patent

Attorney Docket: 892,280-210

·(Formerly 342312004300)

AMENDMENTS TO THE SPECIFICATION

Please replace paragraph [0046] beginning on page 12, line 7 with the following.

[0046] Dalbavancin is thought to inhibit the biosynthesis of the bacterial cell wall by binding to D-alanyl-D-alanine-terminating precursors of peptidoglycans. Dimeric or higher order multimers of dalbavancin may possess further antibacterial properties by interaction of the lipophilic side chains with the cytoplasmic membrane of bacteria. See, for example, Malabarba and Ciabatti, et al. (2001) Current Medicinal Chemistry 8:1759-1773. A further elaboration on dalbavancin multimers may be found in U.S. Serial No. 10/714.166, entitled "DALBAVANCIN COMPOSITIONS FOR TREATMENT OF BACTERIAL

INFECTIONS," filed on November 14, 2003, as Attorney Docket No. 34231-20052,00, the

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disclosure of which is hereby incorporated by reference in its entirety.

Please replace paragraph [0071] beginning on page 17, line 26 with the following.

[0071] Administration and delivery of the drug to the patient, e.g., intravenously, can be done at a controlled rate, so that the concentration in the blood does not increase too quickly or cause precipitation to occur. In some embodiments, dalbavancin is administered at an appropriate rate such that the drug forms a complex with endogenous protein(s) in the bloodstream. Without intending to be bound to a particular theory, it is believed that endogenous protein, such as human serum albumin, can form a complex in vivo with one or two molecules of dalbavancin homolog monomers. When a sufficient amount of dalbavancin is present, it is believed that up to two molecules of dalbavancin homolog will bind to the endogenous protein and it is further

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